CERTIFICATION

SDG No:

1701477C

Laboratory:

Eurofins, Folson, CA

Site:

BMSMC

Matrix:

Air

SUMMARY:

Air samples (Table 1) were collected on the BMSMC facility. The BMSMC facility is located in Humacao, PR. Samples were taken January 26 and 29, 2017 and were analyzed in Eurofins Laboratory of Folson, California that reported the data under SDG No.: 1701477C. The sample results were assessed according to USEPA data validation guidance documents in the following order of precedence: QC criteria from "Compendium Method TO-15. Determination of Volatile Organic Compounds (VOCs) In In Specially-Prepared Canisters and Analyzed Chromatography/Mass Spectrometry (GC/MS), January, 1999"; USEPA Hazardous Waste Support Branch. Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #6. June, 2014). The analyses performed are shown in Table 1. Individual data review worksheets are enclosed for each target analyte group. The data sample summary form shows analytes results that were qualified.

In summary, the results are valid and can be used for decision making purposes.

Table 1. Samples analyzed and analysis performed

SAMPLE ID	SAMPLE DESCRIPTION	MATRIX	ANALYSIS PERFORMED
1701477C-01A	B18SS-2-012617	Air	Methanol
1701477C-02A	B18SS-3-012617	Air	Methanol
1701477C-03A	B18SS-4-012617	Air	Methanol
1701477C-04A	B18SS-5-012617	Air	Methanol
1701477C-05A	B13SS-1-012917	Air	Methanol
1701477C-06A	B13SS-2-012917	Air	Methanol
1701477C-07A	B13SS-2DUP-012917	Air	Methanol
1701477C-08A	B13SS-3-012917	Air	Methanol
1701477C-09A	B15SS-1-012917	Air	Methanol
1701477C-10A	B15SS-1DUP-012917	Air	Methanol
1701477C-11A	B13IA-1-012817	Air	Methanol
1701477C-12A	B13IA-2-012817	Air	Methanol
1701477C-13A	B13IA-2DUP-012817	Air	Methanol
1701477C-14A	B13IA-3-012817	Air	Methanol
1701477C-15A	B18IA-5-012817	Air	Methanol
1701477C-16A	B15IA-1-012817	Air	Methanol
1701477C-17A	B15IA-1DUP-012817	Air	Methanol
1701477C-18A	B1315AA-012817	Air	Methanol

Reviewer Name:

Rafael Infante

Chemist License 1888

Signature:

Date:

March 18, 2017

A 1617194



Client Sample ID: B18SS-2-012617 Lab ID#: 1701477C-01A

EPA METHOD TO-15 GC/MS

2.38	Date of Collection: 1/26/1 Date of Analysis: 2/2/17 0			
Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)	
120	Not Detected	160	Not Detected	
	Rpt. Limit (ppbv)	Rpt. Limit Amount (ppbv) (ppbv)	Rpt. Limit Amount Rpt. Limit (ppbv) (ppbv) (ug/m3)	

Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	101	70-130
Toluene-d8	101	70-130
4-Bromofluorobenzene	99	70-130





Client Sample ID: B185SS-3-012617 Lab ID#: 1701477C-02A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020213 2.42	Date of Collection: 1/26/17 6 Date of Analysis: 2/2/17 06:2		
Compound	Rpt. Limit	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	120	360	160	470

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	98	70-130	
Toluene-d8	103	70-130	
4-Bromofluorobenzene	100	70-130	





Client Sample ID: B18SS-4-012617 Lab ID#: 1701477C-03A

EPA METHOD TO-15 GC/MS

File Name:	j020214		17 5:07:00 PM	
Dil. Factor:	2.82		06:50 PM	
Compound	Rpt. Limit	Amount	Rpt. Limit	Amount
	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	140	160	180	220

Surrogates	%Recovery	Method Limits
1,2-Dichloroethane-d4	98	70-130
Toluene-d8	103	70-130
4-Bromofluorobenzene	105	70-130





Client Sample ID: B18SS-5-012617 Lab ID#: 1701477C-04A

EPA METHOD TO-15 GC/MS

File Name:	j020215	Date of Collection: 1/26/17 5:58:00		
Dil. Factor:	2.52	Date of Analysis: 2/2/17 07:13 PM		
Compound	Rpt. Limit	Amount	Rpt. Limit	Amount
	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	130	140	160	190

		Method
Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	99	70-130
Toluene-d8	104	70-130
4-Bromofluorobenzene	104	70-130





Client Sample ID: B13SS-1-012917 Lab ID#: 1701477C-05A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020216 2.47	Date of Collection: 1/29/17 3:11:00 PM Date of Analysis: 2/2/17 07:36 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	120	190	160	250

		wethod	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	101	70-130	
Toluene-d8	102	70-130	
4-Bromofluorobenzene	104	70-130	





Client Sample ID: B13SS-2-012917 Lab ID#: 1701477C-06A

EPA METHOD TO-15 GC/MS

File Name:	j020217	Date of Collection: 1/29/17 3:55:00 PM		
Dil. Factor:	2.58	Date of Analysis: 2/2/17 07:59 PM		
Compound	Rpt. Limit	Amount	Rpt. Limit	Amount
	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	130	270	170	350

Surrogates	%Recovery	Method Limits
Surrogates	MRECOVERY	Littits
1,2-Dichloroethane-d4	101	70-130
Toluene-d8	103	70-130
4-Bromofluorobenzene	102	70-130





Client Sample ID: B13SS-2DUP-012917 Lab ID#: 1701477C-07A

EPA METHOD TO-15 GC/MS

File Name:	j020218	Date of Collection: 1/29/17 3:55:00 PM		
Dil. Factor:	2.46	Date of Analysis: 2/2/17 08:23 PM		
Compound	Rpt. Limit	Amount	Rpt, Limit	Amount
	(ppby)	(ppbv)	(ug/m3)	(ug/m3)
Compound	(bhna)	(bbns)	(ugmo)	(ug/mə/
Methanol	120	300	160	390

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichtoroethane-d4	97	70-130	
Toluene-d8	105	70-130	
4-Bromofluorobenzene	105	70-130	





Methanol

Air Toxics

Client Sample ID: B13SS-3-012917 Lab ID#: 1701477C-08A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020219 2.58	Date of Collection: 1/29/17 3:36:00 PM Date of Analysis: 2/2/17 08:46 PM		
	Rpt. Limit	Amount	Rpt. Limit	Amount
Compound	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)

130

Container Type: 1 Liter Summa Canister (100% Certified)

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	99	70-130	
Toluene-d8	104	70-130	
4-Bromofluorobenzene	101	70-130	

550



170

720



Client Sample ID: B15SS-1-012917 Lab ID#: 1701477C-09A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020220 2.38	Date of Collection: 1/29/17 5:55:00 P Date of Analysis: 2/2/17 09:10 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	120	230	160	310

Surrogates	%Recovery	Method Limits
1,2-Dichloroethane-d4	101	70-130
Toluene-d8	103	70-130
4-Bromofluorobenzene	102	70-130





Client Sample ID: B15SS-1DUP-012917

Lab ID#: 1701477C-10A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020221 2.47	Date of Collection: 1/29/17 5:55:00 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	120	170	160	220

Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	100	70-130
Toluene-d8	102	70-130
4-Bromofluorobenzene	103	70-130





Client Sample ID: B13IA-1-012817 Lab ID#: 1701477C-11A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020204 1.64	Date of Collection: 1/29/17 2:45:00 PM Date of Analysis: 2/2/17 12:01 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	82	Not Detected	110	Not Detected

	Method	
%Recovery	Limits	
97	70-130	
101	70-130	
101	70-130	
	97 101	





Client Sample ID: B13IA-2-012817 Lab ID#: 1701477C-12A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020205 1.61	Date of Collection: 1/29/17 3:08:00 PM Date of Analysis: 2/2/17 12:24 PM		
	Rpt. Limit	Amount	Rpt. Limit	Amount
Compound	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	80	Not Detected	100	Not Detected

Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	97	70-130
Toluene-d8	102	70-130
4-Bromofluorobenzene	100	70-130





Client Sample ID: B13IA-2DUP-012817

Lab ID#: 1701477C-13A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020206 1.55	Date of Collection: 1/29/17 3:10:00 PM Date of Analysis: 2/2/17 12:47 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	78	Not Detected	100	Not Detected

		Wethod	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	97	70-130	
Toluene-d8	102	70-130	
4-Bromofluorobenzene	103	70-130	





Client Sample ID: B13IA-3-012817 Lab ID#: 1701477C-14A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020207 1.64	Date of Collection: 1/29/17 2:52:00 PI Date of Analysis: 2/2/17 01:10 PM		
Compound	Rpt. Limit	Amount (ppby)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Methanol	82	Not Detected	110	Not Detected

		Method
Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	100	70-130
Toluene-d8	103	70-130
4-Bromofluorobenzene	103	70-130





Client Sample ID: B18IA-5-012817 Lab ID#: 1701477C-15A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020208 1.64	Date of Collection: 1/29/17 4:25:00 PI Date of Analysis: 2/2/17 01:34 PM		
	Rpt. Limit	Amount	Rpt. Limit	Amount
Compound	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	82	82	110	110

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	97	70-130	
Toluene-d8	102	70-130	
4-Bromofluorobenzene	103	70-130	





Client Sample ID: B15IA-1-012817 Lab ID#: 1701477C-16A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020209 1.61		Date of Collection: 1/29/17 3:44:00 PM Date of Analysis: 2/2/17 01:57 PM		
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)	
Methanol	80	Not Detected	100	Not Detected	
Container Type: 6 Liter Sumn	na Canister (100% Certifie	d)			
				Method	
Surrogates		%Recovery		Limits	
1,2-Dichloroethane-d4		97		70-130	
Toluene-d8		103		70-130	
4-Bromofluorobenzene		102		70-130	



Client Sample ID: B15IA-1DUP-012817 Lab ID#: 1701477C-17A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020210 1.55	Date of Collection: 1/29/17 3:44:00 PM Date of Analysis: 2/2/17 02:20 PM		
	Rpt. Limit	Amount	Rpt. Limit	Amount
Compound	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)
Methanol	78	Not Detected	100	Not Detected

Surrogates	%Recovery	Method Limits
1,2-Dichloroethane-d4	98	70-130
Toluene-d8	104	70-130
4-Bromofluorobenzene	101	70-130





Methanol

Air Toxics

Client Sample ID: B1315AA-012817

Lab ID#: 1701477C-18A

EPA METHOD TO-15 GC/MS

File Name: Dil. Factor:	j020211 1.71		te of Collection: 1/29 te of Analysis: 2/2/17	
Compound	Rpt. Limit	Amount	Rpt. Limit	Amount
	(ppbv)	(ppbv)	(ug/m3)	(ug/m3)

Not Detected

86

Container Type: 6 Liter Summa Canister (100% Certified)

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	99	70-130	
Toluene-d8	101	70-130	
4-Bromofluorobenzene	100	70-130	



110

Not Detected



Sample Transportation Notice
Relinquishing signature on this document indicates that sample is being shipped in compliance with all applicable local, State, Federal, national, and international laws, regulations and ordinances of any kind. Air Toxics Limited assumes no liability with respect to the collection, handling or shipping of these samples. Relinquishing signature also indicates agreement to hold harmless, defend, and indemnity Air Toxics Limited against any claim, demand, or action, of any kind, related to the collection, handling, or shipping of samples. D.O.T. Hotline (800) 467-4922

FOLSOM, CA 95630-4719 (916) 985-1000 FAX (916) 985-1020 180 BLUE RAVINE ROAD, SUITE B

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Form 1293 rev.11

21



Chain-of-Custody Record

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EXECUTIVE NARRATIVE

SDG No: 1701477C Laboratory: Eurofins, Folson, CA

Analysis: TO-15 Number of Samples: 18

Location:

SUMMARY: Eighteen (18) samples were analyzed for methanol in ambient air following

Compendium Method TO-15. The sample results were assessed according to USEPA data validation guidance documents in the following order of precedence: QC criteria from "Compendium Method TO-15. Determination of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters and Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999"; USEPA Hazardous Waste Support Branch. Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #6. June, 2014). The QC criteria and data validation actions listed on the data review worksheets are from the primary

guidance document, unless otherwise noted.

Results are valid and can be used for decision making purposes.

Critical issues: None Major: None Minor: None

Critical findings: None Major findings: None

Minor findings: 1. Field duplicates analyzed with this data package. RPD within laboratory and generally

acceptable control limits except for the cases described in the Data Review Worksheet. No action taken, RPD within the method performance criteria and the sample and duplicate

concentration < 5 x SQL.

COMMENTS: Results are valid and can be used for decision making purposes.

Reviewers Name: Rafael Infante

Chemist License 1888

Rafuel Defaut

Signature:

Date: March 16, 2017

METHANOL DATA SAMPLE SUMMARY

METHOD: TO-15

	ı	METHANO	L - TO-15				
Sample ID	Date	Results	Units	Dilution Factor	Lab Flag	Validation	Reportable
1701477C-01A	1/26/2017	160	ug/m³	2.38	-	U	Yes
1701477C-02A	1/26/2017	470	ug/m³	2.42	-	-	Yes
1701477C-03A	1/26/2017	220	ug/m³	2.82	-	-	Yes
1701477C-04A	1/26/2017	190	ug/m³	2.52	-	-	Yes
1701477C-05A	1/29/2017	250	ug/m³	2.47	-	-	Yes
1701477C-06A	1/29/2017	350	ug/m³	2.58	-	-	Yes
1701477C-07A	1/29/2017	390	ug/m³	2.46	-	-	Yes
1701477C-08A	1/29/2017	720	ug/m³	2.58	-	-	Yes
1701477C-09A	1/29/2017	310	ug/m³	2.38	-	-	Yes
1701477C-10A	1/29/2017	220	ug/m³	2.47	-	-	Yes
1701477C-11A	1/29/2017	110	ug/m³	1.64	-	U	Yes
1701477C-12A	1/29/2017	100	ug/m³	1.61	-	U	Yes
1701477C-13A	1/29/2017	100	ug/m³	1.55	-	U	Yes
1701477C-14A	1/29/2017	110	ug/m³	1.64	-	U	Yes
1701477C-15A	1/29/2017	110	ug/m³	1.64	-	-	Yes
1701477C-16A	1/29/2017	100	ug/m³	1.61	-	U	Yes
1701477C-17A	1/29/2017	100	ug/m³	1.55	-	U	Yes
1701477C-18A	1/29/2017	110	ug/m³	1.71	-	U	Yes

	Project Number:1701477C Date:01/26_&_29/17
REVIEW OF VOLATILE ORGATHE following guidelines for evaluating volatile organics was actions. This document will assist the reviewer in using production and in better serving the needs of the data users. The JSEPA data validation guidance documents in the follow Compendium Method TO-15. Determination of Volatile Org Specially-Prepared Canisters and Analyzed By Gas Chalanuary, 1999"; USEPA Hazardous Waste Support Brance Analysis of Ambient Air in Canisters by Method TO-15, (SOP criteria and data validation actions listed on the data review document, unless otherwise noted. The hardcopied (laboratory name) _Eurofins on the quality control and performance data summarized. The data	ere created to delineate required validation of pressional judgment to make more informed the sample results were assessed according to ing order of precedence: QC criteria from anic Compounds (VOCs) In Air Collected Informatography/Mass Spectrometry (GC/MS) h. Validating Air Samples. Volatile Organic # HW-31. Revision #6. June, 2014). The QC w worksheets are from the primary guidance data package received has been reviewed and
_ab. Project/SDG No.:1701477C No. of Samples:18	Sample matrix:Air
Trip blank No.: Field blank No.: Equipment blank No.: Field duplicate No.:_1701477C-06A/1701477C-07A;_17 1701477C-12A/1701477C-13A;_	01477C-09A/1701477C-10A
X Data CompletenessX Sampling Integrity/PreservationX_ GC/MS TuningX_ Internal Standard PerformanceX BlanksX_ Surrogate Recoveries	X Laboratory Control SpikesX Field DuplicatesX CalibrationsX Compound IdentificationsX Compound QuantitationX Quantitation Limits
Overall Comments:VOCs_(methanol)_by_metho	od_TO-15
Definition of Qualifiers: J- Estimated results	

- Compound not detected Rejected data Estimated nondetect U-
- R-
- UJ-

Reviewer:	Rafael defaut	
Date:	03/16/17	

DATA COMPLETENESS

MISSING INFORMATION	DATE LAB. CONTACTED	DATE RECEIVED
		

All criteria were met _	_X
Criteria were not met	
and/or see below	_

SAMPLE INTEGRITY AND PRESERVATION

Canister used for sampling of the ambient air must be demonstrated clean, and leak free prior to sample collection. Cleanliness is demonstrated by the analysis of an individual canister or analysis of a representative canister, if only batch cleaning was required. Leak proof testing is performed on individual canisters. Canisters are used in conjunction with gauges, valves and flow controllers. Therefore, canister should be demonstrated clean and leak free inclusive of these components as appropriate.

a. Leak proof test:

Was the pressure of each canister measured before shipping?

Was the pressure of each canister measured before sampling?

Did the canister hold vacuum/pressure within +/- 2 psi from the date shipped to the sampling date?

Yes or No

Yes or No

Note:

1. The laboratory should be notified if the difference between the laboratory and field pressure is greater than 2 psi.

Actions:

Actions for use of canisters with failing leak test criteria are indicated in Table 1 below.

Table 1. Canister Leak test Actions for TO-15 Analysis*

	Difference in		Action
Matrix	initial and 24 hour pressure (psi) Criteria	Detected Associated Compounds	Non-Detected Associated Compounds
Air	≤ 5	No	qualification
Air	> 5	J	UJ or R

^{*}Excessive time period (> 3months) elapsed between leak test and actual use should be considered in evaluation of canister integrity.

b. Cleanliness

Integrity of the canister used for sampling of air for analysis should be maintained at all times including time of shipment to the field, sampling, shipping back to the laboratory and time of analysis. Analytical results of canister cleaning verification must be taken into account in the validation of sample results.

Does the canister meet the cleanliness criteria?

Yes or No

Is the canister verification included in the data package?

Yes or No

Actions:

Canister contamination actions are stated in Table 2 below.

Note: Laboratory stated that the SUMMA canisters employed were 100 % certified.

Table 2. Canister Contamination Actions for TO-15 Analyses

Contamination Type/level	Canister Cleaning Result	Sample Result	Action for Samples
	Detects	Analytes found in clean canister analysis are non- detects	No qualification required
Clean Canister	<crql< td=""><td>< CRQL</td><td>Report CRQL value with a U</td></crql<>	< CRQL	Report CRQL value with a U
		\geq CRQL and \leq 2x the CRQL	Report concentration of sample with a U
		\geq 2x the CRQL	No qualification required
analysis	> CRQL	< CRQL	Report CRQL value with a U
anarysis		≥ CRQL and ≤ clean canister value	Report clean canister value with a U
		≥ CRQL and > clean canister value	No qualification required
	= CRQL	≤CRQL	Report CRQL value with a U
	CRQL	> CRQL	No qualification required

c. Holding time and sample integrity

SUMMA canisters are to minimize sample charges or loss for majority of the analyte. Sample integrity is maintained by ensuring the system is closed tight and canister pressure from the time of sampling to the time of analysis is maintained within a difference allowable due to temperature change.

Was the canister pressure measured at the conclusion of the sampling period?

Yes or No

Was the canister pressure measured upon arrival to the laboratory? Yes or No Was the canister pressure difference between sampling and analysis less than 5 psi? Yes or No

Actions:

Qualify sample results using technical holding time information as stated in Table 3.

Pressure difference between sampling and analysis should be less than 5 psi. Qualify samples as per Table 3 requirements.

Table 3. Holding Time Actions for TO-15 Volatile Analyses

	Preserved			Action
Matrix	(Pressure difference between sampling and analysis ≤ 5psi)	Criteria	Detected Associated Compounds	Non-Detected Associated Compounds
Air	Yes	< 30 days	N	No qualification
All	Yes	>30 days	J	UJ
Air	No	< 30 days	J	UJ
Air	No	>30 days	J	R

Complete table for all samples and note the integrity and/or preservation not within criteria

SAMPLE ID	DATE SAMPLED	DATE ANALYZED	Pressure difference < 5 psi	ACTION
		commended method ence < 5 psi between		umma canisters received lysis.

The following pressure conversion is used, if necessary

			PRESSURE	CONVERS	ON TABLE			
PSI	ATM	kgf/cm²	in.H₂O	mmHg	in.Hg	Kpa	Bar	mm H₂O
1	0.068046	0.070307	27.7276	51.715	2.03602	6.895	0.6895	704.28104
14.696	1	1.0332	407.484	760	29.921	101.325	1.01325	10350.0936
14.2233	0.96784	1	394.38	735.559	28.959	98.096	0.98067	10000
0.036092	0.002454	0.00253	1	1.8651	0.07343	0.249	0.00249	25.4
0.019336	0.001315	0.001359	0.53616	1	0.03937	0.1333	0.001333	13.618464
0.491154	0.0033421	0.03453	13.6185	25.4	1	3.3864	0.033864	345.9099
0.145	0.00987	0.010197	4.0186	7.5006	0.2953	1	0.01	102.07244
14.5038	0.98692	1.01972	402.156	750.062	29.53	100	1	10214.7624

All criteria were metX	
Criteria were not met see below	

GC/MS TUNING

The assessment of the tuning results is to determine if the sample instrumentation is within the standard tuning QC limits

Gas Chromatograph/Mass Spectrometer (GC/MS) Instrument Performance Check

Action:

NOTES: This requirement does not apply when samples are analyzed by the Selected Ion Monitoring (SIM) technique.

All mass spectrometer instrument conditions must be identical to those used during the sample analysis. Background subtraction actions resulting in spectral distortions for the sole purpose of meeting the method specifications are contrary to the Quality Assurance (QA) objectives, and are therefore unacceptable.

NOTES: No data should be qualified based on BFB or DFTTP failure. Instances of this should be noted in the narrative.

All ion abundance ratios must be normalized to m/z 95, the nominal base peak, even though the ion abundance of m/z 174 may be up to 120% that of m/z 95.

- 1. If samples are analyzed without a preceding valid instrument performance check, qualify all data in those samples as unusable (R).
- 2. If the laboratory has made minor transcription errors which do not significantly affect the data, the data reviewer should make the necessary corrections on a copy of the form.
- 3. If the laboratory has failed to provide the correct forms or has made significant transcription or calculation errors, the Region's designated representative should contact the laboratory and request corrected data. If the information is not available, the reviewer must use professional judgment to assess the data and notify the Project Officer (PO).
- 4. If ion abundance criteria are not met, professional judgment may be applied to determine to what extent the data may be utilized. When applying professional judgment to this topic, the most important factors to consider are the empirical results that are relatively insensitive to location on the chromatographic profile and the type of instrumentation. Therefore, the critical ion abundance criteria for BFB are the m/z 95/96, 174/175, 174/176, and 176/177 ratios. The relative abundances of m/z 50 and 75 are of lower importance. This issue is more critical for Tentatively Identified Compounds (TICs) than for target analytes.
- 5. Note, in the Data Review Narrative, decisions to use analytical data associated with BFB instrument performance check failures (not meeting contract requirements).
- 6. If the reviewer has reason to believe that instrument performance check criteria were achieved using techniques other than those described in the Compendium method TO-15 entitled "Determination Of Volatile Organic Compounds(VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry(GC/MS)", section 10.4, obtain additional information on the instrument performance checks. If the techniques employed are found to be at variance with the contract requirements, the performance and procedures of the laboratory may merit evaluation.
- 7. Use professional judgment to determine whether associated data should be qualified based on the spectrum of the mass calibration compound.

List	the	samples	affected:
If no, use p qualified or i	professional judgment to determin rejected.	e whether the associated data	should be accepted,
XBFE	3 tuning was performed for every 2	24 hours of sample analysis.	
XThe	BFB performance results were re	eviewed and found to be within the	ne specified criteria.

If mass calibration is in error, all associated data are rejected.

All criteria were met	X
Criteria were not met	
and/or see below	

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

Date of initial calibration:	02/02/2017	
Date of initial calibration verificatio	n:	
Dates of continuing calibration:	02/02/2017	
Instrument ID numbers:	MSD-J_	
Matrix/Level:	Air/low	

DATE	LAB FILE ID#	CRITERIA OUT	COMPOUND	SAMPLES
		RFs, %RSD,_%D, r		AFFECTED
Initial and c	ontinuing calibrati	ons meet method spe	ecific requirements. Initial	calibration retention
times meet r	nethod specific re	quirements. One point	calibration curve performe	ed.

Note:

The following criteria apply:

Table 5. Initial Calibration Actions for TO-15 Analyses

	Action		
Criteria for TO-15 Analysis	Detected Associated Compounds	Non-Detected Associated Compounds	
RRF < 0.010 (poor response volatile target compounds, Table 4) RRF < 0.050 (all other volatile target compounds)	J (based on mass spectral R identification)		
RRF > 0.010 (poor response volatile target compounds, Table 4) RRF > 0.050 (all other volatile target compounds)	No qualification		
% RSD > 40.0 or < -40.0 (poor response volatile target compounds, Table 4) % RSD > 30.0 or < -30.0 (all other Volatile target compounds)	No qualification		
% RSD < 40.0 and > -40.0 (poor response volatile target compounds, Table 4) % RSD < 30.0 and > -30.0 (all other volatile target compounds)	J Use profession judgment		

Table 6. Continuing Calibration Verification (CCV) Actions for TO-15 Analyses

	Action		
Criteria for CCV	Detected Associated Compounds	Non-Detected Associated Compounds	
RRF < 0.010 (poor response volatile target compounds, Table 4) RRF < 0.050 (all other volatile target compounds)	J (based on mass spectral identification)	R	
RRF > 0.010 (poor response volatile target compounds, Table 4) RRF > 0.050 (all other volatile target compounds)	No qualification		
%D > 40.0 or < -40.0 (poor response volatile target compounds, Table 4) %D > 30.0 or < -30.0 (all other Volatile target compounds)	J UJ		
%D < 40.0 and > -40.0 (poor response volatile target compounds, Table 4) %D < 30.0 and > -30.0 (all other volatile target compounds)	No qualification		

If the % D for daily calibration exceeds -90, use professional judgment to see if non-detects nee to be qualified as unusable "R"

Note: Methanol is not a poor response compound; the regular calibration/calibration verification criteria are employed.

A separate worksheet should be filled for each initial curve

Table 4. TO 15 Volatile Compounds List*

Compound	CAS Number	Synonyms	
Acetone	67-64-1	Dimethyl ketone; Dimethylformaldehyde; 2-Propanone	
Allyl chloride	107-05-1	3-Chloropropene; 3-Chloroprene	
Benzene	71-43-2	Benzol; Benzine	
Benzyl chloride	100-44-7	Chloromethylbenzene; alpha-Chlorotoluene	
Bromodichloromethane	75-27-4	Monobromodichloromethane; Methane-bromodichloro	
Bromoethene	593-60-2	Vinyl bromide; Monobromoethene	
Bromoform	75-25-2	Tribromoethane	
Bromomethane	74-83-9	Methyl bromide; Monobromomethane	
1,3-Butadiene	106-99-0	Biethylene; Erythrene; Pyrrolyene	
Carbon disulfide	75-15-0	Carbon bisulfide; Carbon sulfide	
Carbon tetrachloride	56-23-5	Carbon tet; Tetrachloromethane	
Chlorobenzene	108-90-7	Monochlorobenzene; Chlorobenzol; Benzene chloride	
Chloroethane	75-00-3	Ethyl chloride; Chlorene; Chloryl	
Chloroethene	75-01-4	Vinyl chloride; Ethylene monochloride	
Chloroform	67-66-3	Trichloromethane; Methyltrichloride; Methane trichloride	
Chloromethane	74-87-3	R40; Methyl chloride; Monochloromethane	
Cyclohexane	110-82-7	Hexamethylene; Hexahydrobenzene; Hexanaphthene	
Dibromochloromethane	124-48-1	Chlorodibromomethane	
1,2-Dibromoethane	106-93-4	EDB; Ethylene dibromide	
1,2-Dichlorobenzene	95-50-1	ODB; Chloroben	
1,3-Dichlorobenzene	541-73-1	meta-Dichlorobenzene; m-Phenylenedichloride	
1,4-Dichlorobenzene	106-46-7	para-Dichlorobenzene; Parazene; Santochlor	
1,1-Dichloroethane	75-34-3	Ethylidene chloride; Ethylidene dichloride	
1,2-Dichloroethane	107-06-2	Ethylene dichloride; Glycol dichloride; 1,2-DCA	
1,1-Dichloroethene	75-35-4	1,1-DCE; Vinylidene chloride	
cis-1,2-Dichloroethylene	156-59-2	cis-1,2-DCE; cis-Acetylene dichloride	
trans-1,2-Dichloroethylene	156-60-5	trans-1,2-DCE; trans-Acetylene dichloride	
1,2-Dichloropropane	78-87-5	Propylene dichloride; Propylene chloride	
cis-1,3-Dichloropropene	10061-01-5	1-Propene,1,3-dichloro-,(z)-; cis-1,3-Dichloro-1-Propene	
trans-1,3-Dichloropropene	10061-02-6	trans-1,3-Dichloro-1-Propene; trans-1,3-Dichloropropylene	
1,4-Dioxane	123-91-1	Diethylene dioxide; Diethylene ether	
Ethyl acetate	141-78-6	Acetic acid ethyl ester; Acetic ether	
Ethylbenzene	100-41-4	Ethylbenzol; Phenylethane	
4-Ethyltoluene	622-96-8	1-Ethyl-4-methyl benzene; p-Methylethylbenzene	
Freon 11 (CCl3F)	75-69-4	Trichlorofluoromethane; Fluorotrichloromethane; Fluorocarbon 11	

Freon 12 (CCl2F2)	75-71-8	Dichlorodifluoromethane; Fluorocarbon 12	
Freon 113 (C2Cl3F3)	76-13-1	1,1,2-Trichloro-1,2,2-trifluoroethane; Fluorocarbon 113; 1,1,2-	
50 09		Trichlorotrifluoroethane	
Freon 114 (C2Cl2F4)	76-14-2	1,2-Dichlorotetrafluoroethane; Halocarbon 114; 1,2-Dichloro-	
		1,1,2,2-tetrafluoroethane	
Heptane	142-82-5	Dipropylmethane; Heptyl hydride	
Hexachlorobutadiene	87-68-3	1,3-Hexachlorobutadiene; Perchlorobutadiene	
Hexane	110-54-3	n-Hexane; Hexyl hydride	
2-Hexanone	591-78-6	Methyl butyl ketone; Butyl methyl ketone; Hexan-2-one	
Isopropyl alcohol	67-63-0	2-Propanol; Isopropanol	
Methylene chloride	75-09-2	Dichloromethane; Methylene dichloride	
Methyl ethyl ketone	78-93-3	MEK; 2-Butanone; Ethyl methyl ketone	
Methyl isobutyl ketone	108-10-1	MIBK; 2-Pentanone; Hexone; Isopropylacetone	
Methyl tert-butyl ether	1634-04-4	MTBE; 2-Methoxy-2-methylpropane; tert-Butyl methyl ether	
Propylene	115-07-1	Propene; Methylethylene	
Styrene	100-42-5	Vinylbenzene; Phenylethylene	
1,1,2,2-Tetrachloroethane	79-34-5	Tetrachloroethane; Acetylene tetrachloride; Bonoform	
Tetrachloroethene	127-18-4	PCE; PERC; Perchloroethylene; Ethylene tetrachloride; Carbon	
		bichloride; Carbon dichloride	
Tetrahydrofuran	109-99-9	Diethylene oxide; Butylene oxide	
Toluene	108-88-3	Toluol; Methylbenzene	
1,2,4-Trichlorobenzene	120-82-1	1,2,4-Trichlorobenzol	
1,1,1-Trichloroethane	71-55-6	Methyl chloroform; Trichloroethane	
1,1,2-Trichloroethane	79-00-5	beta-Trichloroethane; Ethane trichloride; Vinyl trichloride	
Trichloroethene	79-01-6	TCE; Acetylene trichloride; Ethinyl trichloride	
1,2,4-Trimethylbenzene	95-63-6	Pseudocumene; Pseudocumol	
1,3,5-Trimethylbenzene	108-67-8	Mesitylene; Trimethylbenzol	
2,2,4-Trimethylpentane	540-84-1	Iso-octane; Isobutyltrimethylmethane	
Vinyl acetate	108-05-4	Acetic acid ethenyl ether; Ethenyl acetate	
p-Xylene	106-42-3	p-Methyltoluene; 1,4-dimethylbenzene	
m-Xylene	108-38-3	m-Methyltoluene; 1,3-dimethylbenzene	
o-Xylene	95-47-6	o-Methyltoluene; 1,2-Dimethylbenzene	

^{*}Laboratories use different sets and subsets of analytes on as needed basis.

NOTES:

Compounds in bold italicized letters may have poor GCMS response. These poor response compounds are evaluated using more relaxed relative response factor criteria as stated below.

All criteria were met _	
Criteria were not met	
and/or see below	_X

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Blanks criteria and appropriate actions

Blank Type	Blank Result	Sample Result	Action for Samples
	Detects	Not detected	No qualification required
		< CRQL*	Report CRQL value with a U
	< CDOL *	\geq CRQL* and \leq 2x	Report concentration of sample
	< CRQL *	the CRQL**	with a "U"
		\geq 2x the CRQL**	No qualification required
M-4-1 C4	> CRQL *	< CRQL*	Report CRQL value with a U
Method, Storage, Field, Trip,		≥ CRQL* and ≤	Report blank value for sample
Instrument***		blank concentration	concentration with a U
msu ument		≥ CRQL* and >	No qualification required
		blank concentration	No quantication required
	= CRQL*	≤CRQL*	Report CRQL value with a U
		> CRQL*	No qualification required
	Gross	Detects	Report blank value for sample
	contamination **	Detects	concentration with a U

Table 7. Blank Actions for TO-15 Analyses

DATE	LAB ID	LEVEL/	COMPOUND	CONCENTRATION
ANALYZED		MATRIX		UNITS
_All_method_b	lank_meet_meth	nod_specific_cr	iteria	

Note: Concentration detected below the reporting limit, results qualified following the table above.

^{* 2}x the CRQL for methylene chloride, 2-butanone and acetone.

^{** 4}x the CRQL for methylene chloride, 2-butanone, and acetone.

^{***} Qualifications based on instrument blank results affect only the sample analyzed immediately after the sample that has target compounds that exceed the calibration range or non-target compounds that exceed 100 μg/L.

Field/Equipment/Trip blank

DATE ANALYZED	LAB ID	LEVEL/ MATRIX	COMPOUND	CONCENTRATION UNITS
No_field/trip/ed	quipment_blank	s_analyzed_wi	th_this_data_package	
Field/trip blank a	actions			

Field or Trip blank when available should be assessed for possible contaminants in the canister used for trip blank. This canister and its analytical results are specific to the trip blank sample **only**. If contaminants are present in the canister used for trip blank, its suitability for use as trip blank can be assessed using the following criteria.

Table 8. Field/Trip Blank suitability based on Canister contamination

Clean canister Result	Field/Trip Blank Result	Action for Field/Trip Blank
Detects	Not detected	No qualification, no action for samples is required
Detects	<pre>< clean canister result or > clean canister result but < 2X the clean canister result</pre>	Report as non-detect "U", invalid as trip blank, no action for samples is required.
	$\geq 2x$ the clean canister result	No qualification, valid trip blank for sample actions.

CONTAMINATION SOURCE/LEVEL	COMPOUND	CONC/UNITS	AL/UNITS	SQL	AFFECTED SAMPLES

All criteria were met _	_X
Criteria were not met	
and/or see below	

ACTION

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

SURROGATE COMPOUND

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery.

Matrix: solid/aqueous

SAMPLE ID

····· —— ·-	000			
	1,2-DICHLOROETHANE- d4	Toluene- d8	4-BFB	
_Surrogate_recov	eries_within_laboratory_contr	ol_limits		
QC Limits* (Air)				
LL to UL	70 to 130	70 to 13	30 70 to 130	

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

QUALITY	%R < 10%	%R = 10% - LL	%R > UL
Positive results	J	J	J
Nondetects results	R	UJ	Accept

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

All criteria were met _	_X	
Criteria were not met		
and/or see below		

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices. LCS concentration should be in the middle of the calibration range and under the same sample conditions.

1. LCS Recoveries Criteria

Table 9. LCS/LCSD Actions for TO-15 Analyses

	A	Action		
Criteria	Detected Associated Compounds	Non-detected Associated Compounds		
Percent recovery Criteria				
%R > Upper Acceptance Limit (>130%)	J No qualification			
%R in the acceptable range, 70-130%	No qu	ualification		
%R < Lower Acceptance Limit (< 70 %)	J	UJ		
%R < 50%	J	R		
Lower Acceptance Limit \leq %R \leq Upper Acceptance Limit	No qualification			
Relative Percent Difference Criteria				
$\% RPD \le 25\%$	No qu	ualification		
% RPD > 25 %	J	UJ		

	LCS ID	COMPOUND	% R	QC LIMIT
		alyzed_in_this_data_package	_%_recoveries_and	d_RPD
_within_labor	atory_control_limi	its		

2. Frequency Criteria:

Note:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

			All criteria were metX Criteria were not met and/or see below
IX.	FIELD/LABOR	ATORY DUPLICATE PRECISION	
	Sample IDs: Sample IDs: Sample IDs: Sample IDs:	1701477C-06A/1701477C-07A 1701477C-09A/1701477C-10A 1701477C-12A/1701477C-13A 1701477C-16A/1701477C-17A	Matrix:_Air Matrix:_Air Matrix:_Air Matrix:_Air

Field duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information. In the absence of QAPP guidance for validated data from field duplicates, the following action will be taken.

Identify which samples within the data package are field duplicates. Estimate the relative percent difference (RPD) between the vales for each compound. Note large RPDs (>50 %) in the narrative. Use professional judgment to qualify data when RPD > 50 %.

COMPOUND	SQL	SAMPLE	DUPLICATE	RPD	ACTION
		CONC.	CONC.		
1701477C-09A/17	01477C-	-10A			
Methanol	160	310	220	34 %	No action, concentration < 5 SQL
					and RPD within method
					performance criteria.

Laboratory/field duplicate analyzed with this data package. RPD within laboratory and method performance criteria for target analytes except for the cases described in this document. Methanol RPD outside the laboratory control limit but within the method performance criteria, no action taken.

Other suggested actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

All criteria were met	Χ
Criteria were not met	
and/or see below	_

X. INTERNAL STANDARD PERFORMANCE

DATE

SAMPLE ID

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard (CCV standard or mid-point from initial calibration).
- * Retention time (RT) within <u>+</u> 20 seconds of the IS area in the associated calibration standard.

Table 10. Internal Standard Actions for TO-15 Analyses

	Action	
Criteria	Detected Associated Compounds*	Non-detected Associated Compounds*
Area counts > 140% of CCV or mid-point standard from initial calibration)	J-	No qualification
Area counts < 60% of CCV or mid-point standard from initial calibration)	J+	R
Area counts \geq 60% but \leq 140% of CCV or mid-point standard from initial calibration)	No qualification	
RT difference > 20.0 seconds between samples CCV or midpoint standard from initial calibration)	R*	
RT difference < 20.0 seconds between samples and CCV or mid-point standard from initial calibration)	No qualification	

^{*} Examine the chromatographic profile for that sample to determine if any false positives or negatives exist. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for that sample fraction. Detects should not need to be qualified as unusable (R) if the mass spectral criteria are met.

IS OUT

RANGE	
_Internal_standard_area_and_retention_times_within_laboratory_control_limits_for_both_sa _and_calibration_standards	mples

IS AREA

ACCEPTABLE ACTION

All criteria were metX_	
Criteria were not met	
and/or see below	

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

1701477C-02A

Methanol RF = 2.76251

[] = (148261)(400)/(145549)(2.76251)

= 147.5 ppbv OK

All criteria were met	X
Criteria were not met	
and/or see below	_

XII. QUANTITATION LIMITS

A. Dilution performed

SAMPLE ID	DILUTION FACTOR	REASONS FOR DILUTION		
Samples diluted by a factor of 2.82 or less except for the followings.				

System Performance

Action:

Use professional judgment to qualify the data if it is determined that system performance has degraded during sample analyses. Note, for Laboratory Project Officer (PO) action, any degradation of system performance which significantly affected the data.

Note:

Overall Assessment of Data

Action:

- 1. Use professional judgment to determine if there is any need to qualify data which were not qualified based on the Quality Control (QC) criteria previously discussed.
- 2. Write a brief narrative to give the user an indication of the analytical limitations of the data. Note, for Laboratory Project Officer (PO) action, any inconsistency of the data with the Sample Delivery Group (SDG) Narrative. If sufficient information on the intended use and required quality of the data is available, the reviewer should include their assessment of the usability of the data within the given context. This may be used as part of a formal Data Quality Assessment (DQA).

Overall assessment of the data: Results are valid; the data can be used for

decision making purposes.